



Billing Code 4410-09-P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances;
Notice of Registration;
Johnson Matthey, Inc.

By Notice dated November 5, 2013, and published in the
Federal Register on November 18, 2013, 78 FR 69133, Johnson
Matthey, Inc., Pharmaceuticals Materials, 900 River Road,
Conshohocken, Pennsylvania 19428, made application by
renewal to the Drug Enforcement Administration (DEA) to be
registered as a bulk manufacturer of the following basic
classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Oxycodone (9143)	II
Diphenoxylate (9170)	II
Hydrocodone (9193)	II
Meperidine (9230)	II

Drug	Schedule
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers.

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Johnson Matthey, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. The DEA has investigated, Johnson Matthey, Inc., to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk

manufacturer of the basic classes of controlled substances listed.

Dated: February 19, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator,
Office of Diversion Control,
Drug Enforcement Administration.

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